Amendments to the Claims

This listing of claims will replace all prior versions and listings of claims in the application.

1-76. (cancelled).

steroid is a powder.

- 77. (currently amended) A method for preparing a sterile pharmaceutical composition of a steroid comprising:
- (i) dissolving a non-sterile steroid in a solvent to yield a solution of the steroid,
- (ii) filtering the solution of (i) to yield a sterile solution,
- (iii) combining the sterile solution of (ii) with sterile water to form a suspension,
- (iv) optionally removing all or part of the solvent from the suspension of (iii),
- (v) treating the sterile suspension of (iii) or (iv) to obtain a <u>suspension with a particle size</u> distribution having a mass median diameter less than 10 μm,
- (vi) under sterile conditions combining the suspension of (v) with a pharmaceutically acceptable carrier to yield a sterile pharmaceutical composition comprising a suspension of the steroid having a mass median diameter less than 10 μm, and (vii) storing the sterile pharmaceutical composition of (vi) in sterile containers.
- 78. (previously presented) The method of claim 77, wherein the non-sterile

79. (previously presented) The method of claim 78, wherein the powder is a micronized powder.

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- 80. (previously presented) The method of claim 77, wherein the steroid is budesonide.
- 81. (cancelled).
- 82. (previously presented) The method of claim 77, wherein the solvent comprises an alcohol.
- 83. (previously presented) The method of claim 77, wherein the solvent comprises a Class 3 solvent.
- 84. (previously presented) The method of claim 77, wherein the solvent comprises a Class 2 solvent.
- 85. (previously presented) The method of claim 77, comprising combining solvent with the steroid at a temperature from 20 °C below the boiling point of the solvent up to its boiling point.

- 86. (previously presented) The method of claim 85, wherein the solvent is at reflux.
- 87. (previously presented) The method of claim 77, comprising removing solvent under reduced pressure.
- 88. (previously presented) The method of claim 77, comprising removing solvent at atmospheric pressure.
- 89. (previously presented) The method of claim 77, comprising filtering the solution through a filter having a pore size of $0.2 \mu m$ or less.
- 90. (previously presented) The method of claim 77, wherein the sterile water contains a surfactant.
- 91. (previously presented) The method of claim 77, comprising treating the suspension to obtain a particle size distribution having a mass median diameter in the range $1-5 \mu m$.
- 92. (previously presented) The method of claim 91, comprising treating the suspension to obtain a particle size distribution having a mass median diameter in the range $2-3 \mu m$.

- 93. (previously presented) The method of claim 77, comprising storing the sterile composition in sterile ampoules.
- 94. (currently amended) A method for preparing a sterile suspension of budesonide, comprising:
- (i) dissolving non-sterile budesonide in a solvent to yield a budesonide solution,
- (ii) filtering the solution of (i) to yield a sterile solution,
- (iii) combining the sterile solution of (ii) with sterile water to form a suspension of budesonide,
- (iv) optionally removing all or part of the solvent from the suspension of (iii),
- (v) treating the sterile suspension of (iii) or (iv) to obtain a <u>suspension with a particle size</u> distribution having a mass median diameter less than 10 μm,
- (vi) under sterile conditions combining the suspension of (v) with a pharmaceutically acceptable carrier to yield a sterile pharmaceutical composition comprising the suspension of budesonide having a mass median diameter less than 10 μm, and (vii) storing the sterile pharmaceutical composition of (vi) in sterile containers.
- 95. (previously presented) The method of claim 94, wherein the solvent comprises an alcohol.
- 96. (previously presented) The method of claim 94, comprising filtering the solution through a filter having a pore size of 0.2 µm or less.

- 97. (previously presented) The method of claim 96, comprising treating the suspension to obtain a particle size distribution having a mass median diameter in the range $1-5 \, \mu m$.
- 98. (previously presented) The method of claim 96, comprising treating the suspension to obtain a particle size distribution having a mass median diameter in the range $2-3 \mu m$.
- 99.-106. (cancelled).